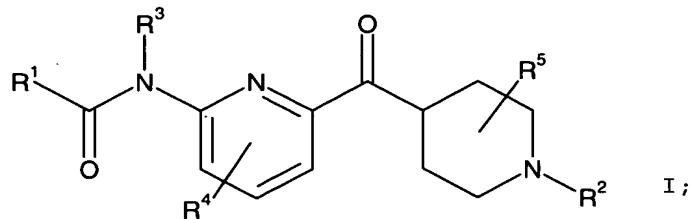


On a new separate last page after the Claims, please insert the following abstract:

ABSTRACT

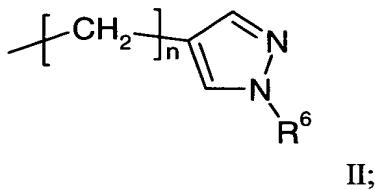
The present invention relates to compounds of formula I:



or pharmaceutically acceptable acid addition salts thereof, where;

R¹ is C₁-C₆ alkyl, substituted C₁-C₆ alkyl, C₃-C₇ cycloalkyl, substituted C₃-C₇ cycloalkyl, C₃-C₇ cycloalkyl-C₁-C₃ alkyl, substituted C₃-C₇ cycloalkyl-C₁-C₃ alkyl, phenyl, substituted phenyl, heterocycle, or substituted heterocycle;

R² is hydrogen, C₁-C₃ alkyl, C₃-C₆ cycloalkyl-C₁-C₃ alkyl, or a group of formula II



II;

R³ is hydrogen or C₁-C₃ alkyl;

R⁴ is hydrogen, halo, or C₁-C₃ alkyl;

R⁵ is hydrogen or C₁-C₃ alkyl;

R⁶ is hydrogen or C₁-C₆ alkyl; and

n is an integer from 1 to 6 inclusively.

The compounds of the present invention are useful for activating 5-HT_{1F} receptors, inhibiting neuronal protein extravasation, and for the treatment or prevention of migraine in a mammal. The present invention also relates to a process for the synthesis of intermediates in the synthesis of compounds of Formula I.